Influence of race/ethnicity on endometriosis

- **43** Fry RP, Crisp AH, Beard RW, McGuigan S. Psychosocial aspects of chronic pelvic pain, with special reference to sexual abuse. A study of 164 women. *Postgrad Med J* 1993;69:566–74.
- **44** Williams C, Long AJ, Noga H, Allaire C, Bedaiwy MA, Lisonkova S, et al. East/South East Asian ethnicity and moderate-to-severe endometriosis. *J Minim Invasive Gynecol* 2019;26:507–15.
- **45** Hudelist G, Fritzer N, Thomas A, Niehues C, Oppelt P, Haas D, et al. Diagnostic delay for endometriosis in Austria and Germany: causes and possible consequences. *Hum Reprod* 2012;27:3412–6.
- **46** Meigs JV. Endometriosis, a possible aetiological factor. *Surg Gynaecol Obstet* 1938;67:253–5.
- 47 Meigs JV. An interest in endometriosis and its consequences: President's address. Am J Obstet Gynecol 1960;79:625–35.
- **48** Blinick G, Merendino VJ. The infrequency of pelvic endometriosis in Negro women. *Am J Surg* 1951;81:635–6.
- 49 Meigs JV. Endometriosis—its significance. Ann Surg 1941;114:866–74.

Disparity in endometriosis diagnoses between racial/ethnic groups

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Bougie and colleagues systematically reviewed the current literature on the association between race/ethnicity and endometriosis diagnosis (Bougie 2019;126:1104–15). BJOGCompared with white women, black women were less likely (OR 0.49, 95% CI 0.29–0.83; n = 16 studies) and Asian women were more likely (OR 1.63, 95% CI 1.03–2.58; n = 10studies) to be diagnosed with endometriosis. The data suggest that compared with white women, Hispanic women were less likely to be diagnosed with endometriosis, with a similar magnitude of effect as black women, but this finding was not stasignificant (OR 0.46, tistically 95% CI 0.14–1.50; n = 5 studies). There are many unanswered questions regarding the interpretation of these findings for research and clinical practice. Ultimately, we do not know whether these associations are artifacts of diagnostic biases or whether there is heterogeneity in endometriosis phenotype or clinical presentation between racial/ethnic groups.

Endometriosis has historically been described as a disease of affluent,

high-achieving women with private health insurance who have delayed marriage and childbearing (Meigs Obstet Gynecol 1953;2:46-53). These early descriptions of endometriosis patients implied a rarity of disease in the 'non-private patient and, therefore by inference (often stated as such) uncommon in black women' (Chatman Am J Obstet Gynecol 1976; 125:987-9), despite early research showing no difference between black and white women and suggesting that pelvic pain in black women was misdiagnosed (Chatman Am J Obstet Gynecol 1976;125:987-9). In 1979, Buttram wrote, 'Typically, our patients with endometriosis appear to have an intense desire to excel. They are usually well-dressed and have trim figures' (Buttram, Jr Fertil Steril 1979;31:117-23). Although overtly sexist, classist and racist descriptions of patients with endometriosis are less common today, these not-so-distant views may still consciously or unconsciously influence clinical care, leading to disparities in endometriosis diagnoses.

Differences in endometriosis diagnoses between black, Asian,

Hispanic, and white women are likely to be synergistic with larger systemic problems related to the difficulty of receiving an endometriosis diagnosis. Indeed, women globally wait, on average, 7 years between the onset of endometriosis symptoms and disease diagnosis, with some women never receiving a formal diagnosis. There are numerous factors that contribute to this diagnostic delay, including inconsistent symptom recognition by both the patient and the provider, and the lack of a non-invasive diagnostic test. When the results of the present study were restricted to women presenting with infertility, disparities for black women attenuated and there was no statistically significant difference endometriosis diagnoses black, white, and Asian patients, suggesting that once patients establish access to clinical care, disparities in diagnosis diminish.

Bougie et al. described substantial heterogeneity ($I^2 > 50\%$) between studies. Although the authors stratified by study year and diagnosis modality, differences in access to

health care across populations are also likely to contribute to this heterogeneity, given the social and economic complexities of diagnosis. Different countries will have different medical referral patterns and insurance systems, as well as different racial and economic biases. Both race and gender have been shown to unconsciously influence referral patterns and clinical care (Schulman et al. *N Engl J Med* 1999;340:618–26). Although future studies investigating the association between race/ethnicity and endometriosis are

warranted, they must be interpreted through the lens of quantifying disparities in access to diagnostic care. We must ensure that women of all racial and ethnic backgrounds are comfortable recognising and describing their symptoms, and that healthcare providers take these symptoms seriously and provide appropriate and timely referral for endometriosis diagnosis and treatment.

Disclosure of interests

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